

Product datasheet for SR413518

OriGene Technologies, Inc.

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Eif4a1 Mouse siRNA Oligo Duplex (Locus ID 13681)

Product data:

Product Type: siRNA Oligo Duplexes

Purity: HPLC purified

Quality Control: Tested by ESI-MS

Sequences: Available with shipment

Stability: One year from date of shipment when stored at -20°C.

of transfections: Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final

conc. 10 nM).

Note: Single siRNA duplex (10nmol) can be ordered.

RefSeq: <u>NM 001159375, NM 144958</u>

UniProt ID: P60843

Synonyms: BM-010; Ddx2a; Eif4

Components: Eif4a1 (Mouse) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 13681)

Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol

Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 ml

Summary: ATP-dependent RNA helicase which is a subunit of the eIF4F complex involved in cap

recognition and is required for mRNA binding to ribosome. In the current model of

translation initiation, eIF4A unwinds RNA secondary structures in the 5'-UTR of mRNAs which

is necessary to allow efficient binding of the small ribosomal subunit, and subsequent

scanning for the initiator codon.[UniProtKB/Swiss-Prot Function]







Performance Guaranteed:

OriGene guarantees that at least two of the three Dicer-Substrate duplexes in the kit will provide at least 70% or more knockdown of the target mRNA when used at 10 nM concentration by quantitative RT-PCR when the TYE-563 fluorescent transfection control duplex (cat# SR30002) indicates that >90% of the cells have been transfected and the HPRT positive control (cat# SR30003) provides 90% knockdown efficiency.

For non-conforming siRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the siRNA kit. To arrange for a free replacement with newly designed duplexes, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled siRNA control (quantitative RT-PCR data required).